

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
(Case No. 99,424-T1)

INFORMATION DISCLOSURE STATEMENT

In the judgment of the undersigned, portions of the listed references may be material to the Examiner's consideration of the presently pending claims. This statement is not a representation that the listed references have effective dates early enough to be "prior art" within the meaning of 35 U.S.C. Section 102 or Section 103.

Applicants do not believe any fee is due with this submission. If this belief be in error and the Patent Office determines that the fee prescribed in the relevant portion of 37 C.F.R. Section 1.97 is applicable, the undersigned attorney by his signature hereby authorizes any such fee to be debited from Deposit Account 13-2490.

U. S. PATENTS

1. Miyashita et al., U.S. Patent No. 5,189,027, issued February 23, 1993
2. Cristalli, U.S. Patent No. 5,593,975, issued January 14, 1997
3. Miyasaka et al., U.S. Patent No. 4,956,345, issued September 11, 1990
4. Kogi et al., U.S. Patent No. 5,270,304, issued December 14, 1993
5. Yamaguchi et al., U.S. Patent No. 5,459,254, issued October 17, 1995
6. Yamada, U.S. Patent No. 5,705,491, issued January 6, 1998
7. Khan et al., U.S. Patent No. 5,770,716, issued June 23, 1998
8. Morozumi et al., U.S. Patent No. 5,939,543, issued August 17, 1999
9. Zablocki et al., U.S. Patent No. 6,214,807, issued April 10, 2001
10. Verani, U.S. Patent No. 6,026,317, issued February 15, 2000

FOREIGN PATENT DOCUMENTS

11. Canadian Patent No. 965,411, published April 1, 1975
12. Japanese Patent No. 5[1993]-9197, published January 19, 1993
13. European Patent No. 0 354,638, published February 14, 1990

OTHER DOCUMENTS

14. Marumoto, et al., "Synthesis and Coronary Vaso-dilating Activity of 2-Substituted Adenosines", *Chem. Pharm. Bull.* 23(4): 759-774 (1975).
15. Marumoto, et al., "Synthesis and Enzymatic Activity of Adenosine 3',5'-Cyclic Phosphate Analogs", *Chem. Pharm. Bull.* 27(4) 900-1003 (1979).
16. Persson, et al., "Synthesis and Antiviral Effects of 2-Heteroaryl Substituted Adenosine and 8-Heteroaryl Substituted Guanosine Derivatives", *Bioorganic & Medicinal Chemistry*, 3:1377-1382 (1995).
17. Mager, et al., "Molecular simulation applied to 2-(N'-alkylidenehydrazino)- and 2-(N'-aralkylidenehydrazino) adenosine A₂ Agonists", *Eur J. Med. Chem*, 30:15-25 (1995).
18. Cristalli et al., "2-Alkynyl Derivatives of Adenosine 5'-N'-ethyluronamide: Selective A₂ Adenosine Receptor Agonists with Potent Inhibitory Activity on Platelet Aggregation", *J. Med. Chem*,

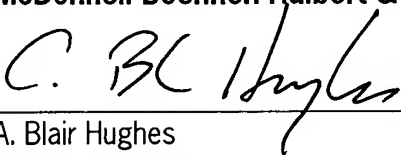
37:1720-1726 (1994).

19. Matsuda, et al., "Nucleosides and Nucleotides. 103. 2-Alkynyladenosines: A Novel Class of Selective Adenosine A₂ Receptor Agonists with Potent Antihypertensive Effects", *J. Med. Chem.* 35:241-252 (1992).

Respectfully submitted,
McDonnell Boehnen Hulbert & Berghoff LLP

Date: March 30, 2004

By:


A. Blair Hughes
Reg. No. 32,901

FORM PTO-1449
(Rev. 2-32)

U.S. Department of Commerce
Patent and Trademark Office

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Use several sheets if necessary)

Atty. Docket No.

99,424-T1

Serial No.

Not Assigned

Applicant:

Jeff Zablocki et al.

Filing Date:

Concurrently
Herewith

Group:

U.S. PATENT DOCUMENTS

Examiner Initial	Document Number	Date	Name	Class	Subclasses	Filing Date if Appropriate
	5,189,027	2/23/93	Miyashita et al.			
	5,593,975	1/14/97	Cristalli			
	4,956,345	9/11/90	Miyasaka et al.			
	5,270,304	12/14/93	Kogi et al.			
	5,459,254	10/17/95	Yamaguchi et al.			
	5,705,491	1/6/98	Yamada			
	5,770,716	6/23/98	Khan et al.			
	5,939,543	8/17/99	Morozumi et al.			
	6,214,807	4/10/01	Zablocki et al.			
	6,026,317	2/15/00	Verani			

FOREIGN PATENT DOCUMENTS

	Document Number	Date	Country	Class	Subclass	Translation	
						Yes	No
	965,411	4/1/75	Canada				
	Hei 5[1993]-9197	1/19/93	Japan				
	0 354 638	2/14/90	Europe				

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc).

Marumoto, et al., "Synthesis and Coronary Vaso-dilating Activity of 2-Substituted Adenosines", *Chem. Pharm. Bull.* 23(4): 759-774 (1975).

EXAMINER

DATE CONSIDERED

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication.

FORM PTO-1449
(Rev. 2-32)

U.S. Department of Commerce
Patent and Trademark Office

Atty. Docket No.

99,424-T11

Serial No.

Not Assigned

**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**
(Use several sheets if necessary)

Applicant:

Jeff Zablocki et al.

Filing Date:

Concurrently
Herewith

Group:

U.S. PATENT DOCUMENTS

Examiner Initial		Document Number	Date	Name	Class	Subclas s	Filing Date if Appropriate

FOREIGN PATENT DOCUMENTS

		Document Number	Date	Country	Class	Subclass	Translation	
							Yes	No

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc).

		Marumoto, et al., "Synthesis and Enzymatic Activity of Adenosine 3'5'-Cyclic Phosphate Analogs", Chem. Pharm. Bull. 27(4) 990-1003 (1979).
		Persson, et al., "Synthesis and Antiviral Effects of 2-Heteroaryl Substituted Adenosine and 8-Heteroaryl Substituted Guanosine Derivatives", <i>Bioorganic & Medicinal Chemistry</i> , 3:1377-1382 (1995).
		Mager, et al., "Molecular simulation applied to 2-(N'alkylidenehydrazino)- and 2-(N'-aralkylidenehydrazino) adenosine A ₂ Agonists", <i>Eur J. Med. Chem.</i> , 30:15-25 (1995).
		Cristalli et al., "2-Alkynyl Derivatives of Adenosine 5'-N'ethyluronamide: Selective A ₂ Adenosine Receptor Agonists with Potent Inhibitory Activity on Platelet Aggregation", <i>J. Med. Chem.</i> , 37:1720-1726 (1994).
		Matsuda, et al., "Nucleosides and Nucleotides. 103. 2-Alkynyladenosines: A Novel Class of Selective Adenosine A ₂ Receptor Agonists with Potent Antihypertensive Effects", <i>J. Med. Chem.</i> 35:241-252 (1992).

EXAMINER

DATE CONSIDERED

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication.